

# Unconventional resistivity at the border of metallic antiferromagnetism in $\text{NiS}_2$

P. G. Niklowitz,\* M. J. Steiner, and G. G. Lonzarich  
*Cavendish Laboratory, University of Cambridge, Madingley Road, Cambridge CB3 0HE, UK*

D. Braithwaite, G. Knebel, and J. Flouquet  
*Département de Recherche Fondamentale sur la Matière Condensée,  
 SPSMS, CEA Grenoble, 38054 Grenoble Cedex 9, France*

J. A. Wilson  
*H. H. Wills Physics Laboratory, University of Bristol, Bristol BS8 1TL, UK*  
 (Dated: February 6, 2008)

We report low-temperature and high-pressure measurements of the electrical resistivity  $\rho(T)$  of the antiferromagnetic compound  $\text{NiS}_2$  in its high-pressure metallic state. The form of  $\rho(T)$  suggests that metallic antiferromagnetism in  $\text{NiS}_2$  is quenched at a critical pressure  $p_c = 76 \pm 5$  kbar. Near  $p_c$  the temperature variation of  $\rho(T)$  is similar to that observed in  $\text{NiS}_{2-x}\text{Se}_x$  near the critical composition  $x = 1$  where the Néel temperature vanishes at ambient pressure. In both cases  $\rho(T)$  varies approximately as  $T^{1.5}$  over a wide range below 100 K. However, on closer analysis the resistivity exponent in  $\text{NiS}_2$  exhibits an undulating variation with temperature not seen in  $\text{NiSe}$  ( $x = 1$ ). This difference in behaviour may be due to the effects of spin-fluctuation scattering of charge carriers on cold and hot spots of the Fermi surface in the presence of quenched disorder, which is higher in  $\text{NiSe}$  than in stoichiometric  $\text{NiS}_2$ .

PACS numbers: 71.27.+a, 71.10.-w, 72.80.Ga, 75.40.-s

Keywords: strongly correlated electron systems, transition metal compounds,  $\text{NiS}_2$ , electronic transport, spin fluctuations, itinerant antiferromagnetism

## I. INTRODUCTION

The electronic properties of metals on the border of magnetic phase transitions at low temperatures are often found to exhibit temperature dependences that differ from the predictions of Fermi liquid theory. Early attempts to explore such non-Fermi liquid behaviour have been based on mean-field treatments of the effects of enhanced magnetic fluctuations, as in the self-consistent renormalization (SCR) model.<sup>1,2,3,4</sup>

In a recent work the prediction of this model for the temperature dependence of the electrical resistivity  $\rho(T)$  was tested in a simple cubic d-metal,  $\text{Ni}_3\text{Al}$ , at high pressures near to the critical pressure where ferromagnetism is suppressed.<sup>5</sup> The  $T^{5/3}$  temperature dependence of the resistivity seen in  $\text{Ni}_3\text{Al}$  and other related systems, where the magnetic correlation wavevector  $\kappa(T)$  is small, appears to be largely consistent with the SCR model.<sup>6</sup> In the idealized limit  $\kappa \rightarrow 0$  at  $T \rightarrow 0$ , the SCR model predicts that in 3D the quasiparticle scattering rate  $\tau_{qp}^{-1}$  varies linearly with the quasiparticle excitation energy, rather than quadratically as in the standard Fermi liquid picture. This form of  $\tau_{qp}^{-1}$  is similar to that of the marginal Fermi liquid model,<sup>7,8,9</sup> which is normally associated with a linear temperature dependence of the resistivity. However, at the border of ferromagnetism the relevant fluctuations responsible for quasiparticle scattering are of long wavelength and thus are ineffective in reducing the current. This leads to a transport relaxation rate  $\tau_{tr}^{-1}$  that differs from  $\tau_{qp}^{-1}$  and varies not as  $T$ , but as  $T^{5/3}$ .

In practice, this form of  $\tau_{tr}^{-1}$  or of the resistivity  $\rho(T)$  can be restricted to a relatively narrow range of temperatures and pressures where  $\kappa(T)$  is small compared with the characteristic wavevector of thermally excited magnetic fluctuations. The SCR model is not restricted to this limit and can in principle include the effects of  $\kappa(T)$  and even determine  $\kappa(T)$  in a self-consistent fashion. We note that the SCR model assumes implicitly that an effective underlying mechanism exists to remove momentum from the electron system. The validity of this assumption has not been clearly confirmed theoretically, but seems to be consistent with experiment in the cases mentioned above. In its conventional form the model does not include the possible effects of inhomogeneities or texture that may arise on the border of first order ferromagnetic transitions as in, e.g.,  $\text{MnSi}$ .<sup>13,14,15</sup> The SCR model is also expected to fail on the border of electron localization, as near a Mott transition or close to local quantum critical points in heavy electron compounds.<sup>16,17</sup>

The applicability of the SCR model has also been questioned for the case of itinerant-electron antiferromagnetism in general, even well away from the border of a Mott transition.<sup>18,19,20</sup> In this paper we present an attempt to test the prediction of the SCR model in a metal on the border of metallic antiferromagnetism in 3D for which  $\rho$  is predicted to vary as  $T^{3/2}$  in the idealized limit  $\kappa \rightarrow 0$ ,  $T \rightarrow 0$ , where  $\kappa$  now stands for the correlation wavevector for the staggered magnetization. The exponent of  $3/2$  is the ratio of the spatial dimension  $d = 3$  and the dynamical exponent  $z = 2$ . This may be contrasted with the corresponding exponent of  $5/3$ , which is the ratio of  $(d + 2) = 5$  and  $z = 3$ , for the border

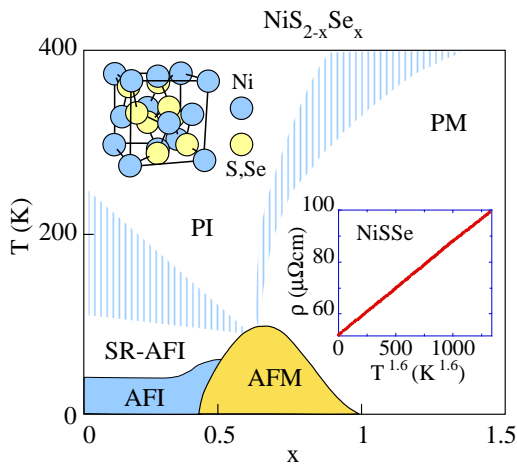


FIG. 1: Temperature-composition phase diagram of  $\text{NiS}_{2-x}\text{Se}_x$ .<sup>10,11</sup> Lower inset shows the temperature dependence of the resistivity of  $\text{NiSSe}$  ( $x = 1$ ), which is just on the border of a metallic antiferromagnetic state at low temperatures.<sup>12</sup> The upper inset is the pyrite crystal structure of  $\text{NiS}_2$  and  $\text{NiSe}_2$ . PI and PM stand for paramagnetic insulator and paramagnetic metal, respectively. AFI and AFM stand for antiferromagnetic insulator and antiferromagnetic metal, respectively. SR-AFI stands for short range antiferromagnetic insulator.<sup>11</sup> The temperature-pressure phase diagram of  $\text{NiS}_2$  is expected to be similar in form to the temperature-composition phase diagram of  $\text{NiS}_{2-x}\text{Se}_x$ . In this study, the AFM boundary of  $\text{NiS}_2$  is found to be at  $p_c = 76 \pm 5$  kbar. Thus,  $\text{NiS}_2$  at  $p_c$  and  $\text{NiSSe}$  at ambient pressure may be expected to be similar except for the level of quenched disorder, which is normally higher in  $\text{NiSSe}$  than in  $\text{NiS}_2$ . The residual resistivity of  $\text{NiSSe}$  is typically an order of magnitude higher than in  $\text{NiS}_2$ .

of metallic ferromagnetism. (The 2 in  $d + 2$  arises from the effect of small-angle scattering that is absent in the case of the staggered magnetization.) This simple model for the scattering from antiferromagnetic fluctuations assumes that the scattering rate can be averaged over the Fermi surface. Within the SCR model this procedure would seem to require the presence of a sufficient level of quenched disorder assumed to have only the simple consequence of inhibiting the short circuiting caused by the carrier on the cold spots of the Fermi surface, i.e., regions far from the hot spots connected by the antiferromagnetic ordering wavevector and thus strongly affected by spin-fluctuation scattering.<sup>18,19,20</sup>

The effect of quenched disorder on the temperature dependence of  $\rho$  in the SCR model shows up particularly clearly in the temperature-dependent resistivity exponent defined as  $n = \partial \ln \Delta\rho / \partial \ln T$ , where  $\Delta\rho = \rho(T) - \rho_0$  and  $\rho_0$  is the residual resistivity, i.e., the resistivity extrapolated to  $T = 0$  K. The resistivity exponent  $n$  may be described in terms of the Fermi wavevector  $k_F$ , the elastic mean free path  $l$  of charge carriers and the reduced temperature  $t = T/T_{sf}$ , where  $T_{sf}$  is a characteristic spin fluctuation temperature.<sup>6</sup> With increasing  $t$ ,  $n$  drops from  $3/2$  towards unity around  $t \approx 1/(k_F l)$ , back

towards a value of the order of  $3/2$  around  $t \approx 1/\sqrt{k_F l}$  and then towards zero for  $t \gg 1/(k_F l)$ .<sup>18,19,20</sup> This undulating behaviour of  $n$  is a dramatic prediction of the model for relatively pure samples that could in principle be tested by studying a series of samples of the same material with different levels of quenched disorder. Rosch<sup>19,20</sup> introduced this model in an effort to understand the temperature dependence of the resistivity exponent measured for the f-electron metal  $\text{CePd}_2\text{Si}_2$ .<sup>21,22</sup> The results were, in this case, not entirely conclusive because of the anisotropic and complex nature of the spin fluctuation spectrum in this material.

Here we present a test of Rosch's model in a d-electron system with a cubic structure and  $T_{sf}$  one to two orders of magnitude greater than in typical heavy f-electron systems. We compare the temperature dependence of  $\rho$  of two closely related systems on the border of antiferromagnetism. The two materials are  $\text{NiS}_2$  near 76 kbar and  $\text{NiS}_{2-x}\text{Se}_x$  for  $x = 1$  where the Néel temperature  $T_N$  vanishes at ambient pressure. Due to random variations in the locations of the S and Se atoms in the lattice, values of  $\rho_0$  found in  $\text{NiSSe}$  are typically one order of magnitude higher than in stoichiometric  $\text{NiS}_2$  compounds.

$\text{NiS}_2$  crystallizes in the cubic pyrite structure and is an antiferromagnetic insulator at ambient pressure at low temperatures.<sup>23,24</sup> It can be metallized via the application of pressure<sup>25,26,27,28</sup> or by Se substitution.<sup>10,12,25,27,28,29,30</sup> The temperature-pressure phase diagram of  $\text{NiS}_2$  and the temperature-composition phase diagram of  $\text{NiS}_{2-x}\text{Se}_x$  (Fig. 1) are expected to be similar. Of particular interest for this paper is the boundary of metallic antiferromagnetism that appears (i) at  $x \cong 1$ , i.e., for  $\text{NiSSe}$ , in temperature-composition phase diagram (Fig. 1) and (ii) at  $p_c$  in the temperature-pressure phase diagram for stoichiometric  $\text{NiS}_2$ . We note that in both cases the quantum critical point for metallic antiferromagnetism arises well beyond the metal insulator transition (see Fig. 1 for the case where composition is the control parameter).

We present a high-pressure study of the temperature dependence of the resistivity of  $\text{NiS}_2$  in the metallic state, which reveals a critical pressure  $p_c \cong 76$  kbar. The results at  $p_c$  are compared with that of  $\text{NiSSe}$  at ambient pressure reported in Ref. 12. In both materials one observes a non-Fermi liquid form of the resistivity that appears in first approximation to be consistent with the predictions of the SRC model (inset of Fig. 1 for  $\text{NiSSe}$ ). However, the resistivity of  $\text{NiS}_2$  near  $p_c$  has an undulating component in its variation with temperature that is not seen in  $\text{NiSSe}$ . This difference in behaviour might arise from the effects of cold and hot spots on the Fermi surface as suggested by the model due to Rosch discussed above.<sup>19,20</sup>

## II. EXPERIMENTAL

Single crystals of  $\text{NiS}_2$  have been grown by chemical vapour transport using iodine as the transport

agent.<sup>25</sup> The residual resistivity ratio ( $\rho(273\text{ K})/\rho_0$ ) above 50 kbar is about 30 for our samples, compared with about 3 for NiSSe.<sup>12</sup> The carrier mean free path of NiS<sub>2</sub> is thus expected to be about an order of magnitude higher than in NiSSe.

Pressure was applied by means of a non-magnetic Bridgman cell using tungsten carbide anvils. The cell used is a scaled down version of that designed by Wittig.<sup>31</sup> The cullet diameter of the anvils was 3.5 mm. The gasket was made of pyrophyllite with a central hole to accommodate the NiS<sub>2</sub> sample as well as a Pb sample and the steatite pressure-transmission medium. After compression, the sample space was reduced to about 1.5 mm in diameter and 0.1 mm in thickness. The pressure was determined from the superconducting transition temperature of the Pb sample.<sup>32</sup> The width of the superconducting transition suggested that the pressure variation over the Pb sample was about 10% of the average pressure.

The resistivity was measured via the 4-terminal ac technique. Four 50  $\mu\text{m}$  Pt leads were passed into the high-pressure region via grooves in the insulating pyrophyllite gasket. The bare wires were rested on top of the samples and pressed onto the sample surface during pressurization to achieve adequate electrical contacts.

The resistivity measurements were carried out with two different and independent systems, a helium circulation cryostat (ILL Orange cryostat; 1.5 K to 300 K) in Grenoble and an adiabatic demagnetization refrigerator (0.04 K to 100 K) in Cambridge. The latter system had two voltage channels, one with a low temperature transformer and the other with a room-temperature transformer. The excitation currents were 1 mA and 160  $\mu\text{A}$  in the Orange cryostat and in the adiabatic demagnetization fridge, respectively. The results obtained with these two experimental systems were consistent with each other where comparisons could be made.

### III. RESULTS

The temperature variations of the resistivity of NiS<sub>2</sub> in the high pressure metallic state above 40 kbar are presented in Figures 2 and 3. (The insulator-to-metal transition (not shown) was observed at around 30 kbar, in agreement with the literature.<sup>25</sup>) Figure 2 shows  $\Delta\rho$  vs  $T$  up to 80 K and Figure 3 is an expanded view of  $\Delta\rho$  vs  $T$  in the range 0.05 K to 2 K. Plots of  $\Delta\rho$  vs  $T^2$  and  $\Delta\rho$  vs  $T^{3/2}$  over different ranges in temperatures are shown, respectively, in Figures 5 and 6.

Below 1 K the resistivity can be described by an equation of the form  $\rho = \rho_0 + AT^2$  over the entire pressure range explored, 43 kbar to 96 kbar. The pressure variations of the fitted values of  $A$  and  $\rho_0$  are given in Figure 4. The  $T^2$  coefficient  $A$  exhibits a peak at  $p_c = 76 \pm 5$  kbar.

Figures 5 and 6 compare  $\Delta\rho$  vs  $T^2$  and  $\Delta\rho$  vs  $T^{3/2}$ , respectively, in three panels each covering different ranges in temperature. Figure 5c, in particular, highlights the

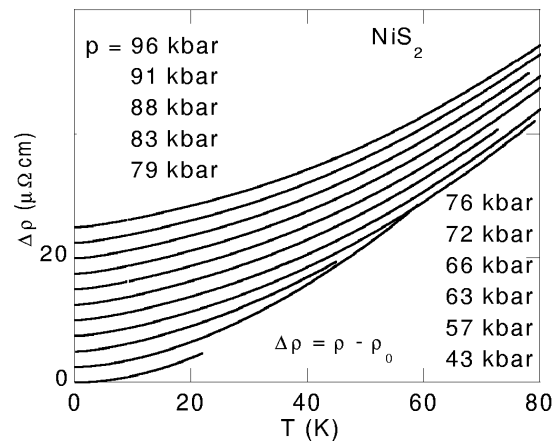


FIG. 2: Temperature dependence of the resistivity in the metallic state of NiS<sub>2</sub> at high pressures.  $\Delta\rho$  is  $\rho(T) - \rho_0$ , where  $\rho_0$  is the residual resistivity extrapolated to  $T = 0$  K. The curves are shifted vertically for clarity.

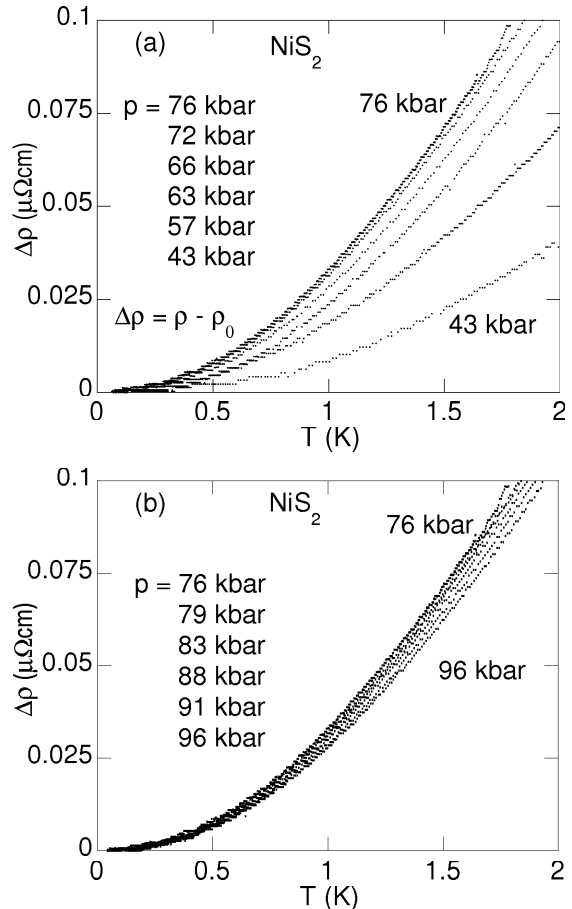


FIG. 3: Low temperature variation of the resistivity in the metallic state of NiS<sub>2</sub> at high pressures.  $\Delta\rho$  grows in strength up to 76 kbar (a) and weakens gradually above 76 kbar (b).

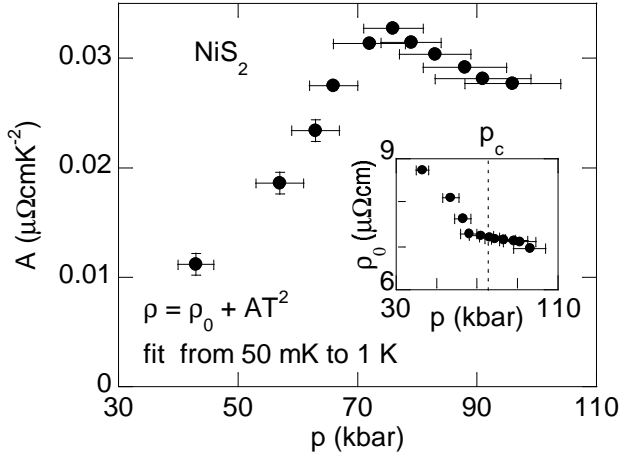


FIG. 4: The  $T^2$  coefficient of the resistivity,  $A$ , and residual resistivity,  $\rho_0$  (inset), of  $\text{NiS}_2$  in the metallic high-pressure state. The parameters  $A$  and  $\rho_0$  are obtained by a fit of the resistivity as indicated in the figure.  $A$  is peaked at  $p_c = 76 \pm 5$  kbar. This marks the boundary of the metallic antiferromagnetic state of  $\text{NiS}_2$ .

$T^2$  variation observed at all pressures with a peak of  $A$  at around  $p_c$  as discussed above. Figures 6a, 6b and 7, on the other hand, suggest that near this pressure  $\Delta\rho$  varies roughly as  $T^{3/2}$  over a decade in temperature above a few K. This is the behaviour predicted by the SCR model for a system on the border of metallic antiferromagnetism as discussed in the introduction. The existence of a  $T^2$  regime even at  $p_c$ , however, suggests that the transition into the antiferromagnetic state below  $p_c$  may not be continuous, i.e., that the antiferromagnetic quantum critical point is not quite reached due to the onset of a first order transition. We note that the weak pressure variation of the Fermi liquid crossover temperature  $T_{FL}$  near  $p_c$  (see Fig. 7 and the caption) would seem to rule out an explanation of the  $T^2$  resistivity in terms of an inhomogeneity in pressure.

The identification of  $p_c$  with the boundary of metallic antiferromagnetism is also suggested by the correspondence of the temperature-pressure and temperature-composition phase diagram of  $\text{NiS}_2$  found in earlier work.<sup>10,12,29,30</sup> This predicted that the critical pressure for the border of metallic antiferromagnetism should be around 60 kbar, which is of the same order of magnitude as  $p_c$  defined above. The identification of  $p_c$  with the antiferromagnetic boundary in the metallic state is tentative and needs confirmation by other measurements and in particular the detection of a signature of  $T_N$  in the resistivity for pressures below  $p_c$ . Here we focus attention mainly on the behaviour of  $\rho$  near  $p_c$  and contrast it with that of  $\text{NiSSe}$  at ambient pressure (inset of Fig. 1).

The non-Fermi liquid behaviour of  $\Delta\rho$  over a wide temperature range near  $p_c$  (low-temperature data at 76 kbar with added 77 kbar data up to 300 K; the high-temperature data is scaled to match the low-temperature data) is highlighted in Figure 8. Panel (a) shows the

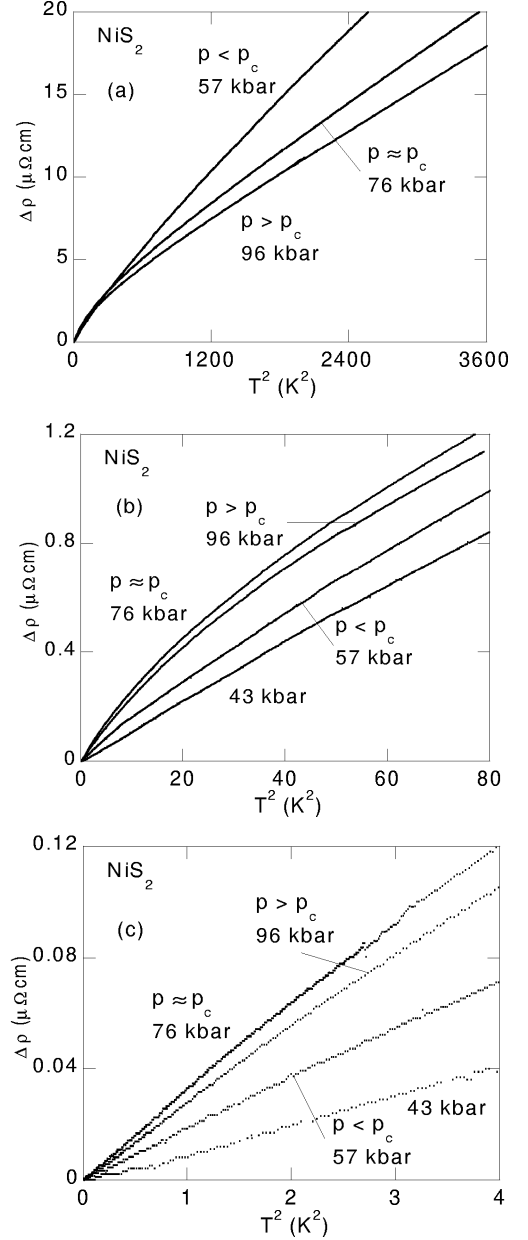


FIG. 5: Resistivity vs  $T^2$  in the high pressure metallic regime of  $\text{NiS}_2$ . The three panels are for different temperature intervals. A quadratic temperature dependence of  $\Delta\rho$  is seen only at the lowest temperatures (below a few K) and the  $T^2$  coefficient of  $\Delta\rho$  is peaked at  $p_c \approx 76$  kbar as shown in Fig. 4.

dramatic upturn of  $\Delta\rho/T^2$ , which in the Fermi liquid regime is expected to saturate to a constant value. Panel (b) is a plot of  $\ln \Delta\rho$  vs  $\ln T$  which shows that the average slope corresponds to a resistivity exponent close to  $3/2$ , as discussed above and as seen in  $\text{NiSSe}$  at ambient pressure. However, in contrast to  $\text{NiSSe}$ , the temperature dependence of the resistivity in  $\text{NiS}_2$  at  $p_c$  exhibits an undulating structure which is evident in Figure 8 and highlighted in the temperature dependence of the resistivity exponent  $n = \partial \ln \Delta\rho / \partial \ln T$  shown in the inset.



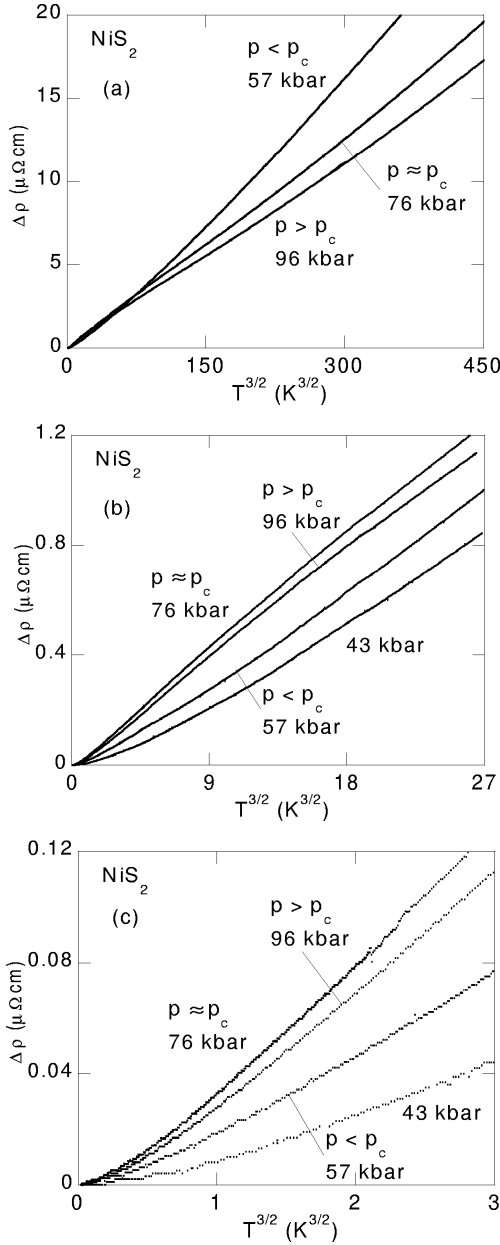


FIG. 6: Resistivity vs  $T^{3/2}$  in the high-pressure metallic regime of  $\text{NiS}_2$ . The three panels are for different temperature intervals. An approximately  $T^{3/2}$  variation of  $\Delta\rho$  is seen around  $p_c \approx 76$  kbar only over a decade in temperature above a few K.  $\Delta\rho$  is quadratic in temperature below a few K at all pressures in the metallic regime studied (see Fig. 5c).

We note that this undulating structure has been observed in several samples of  $\text{NiS}_2$  and in two independent measurement systems.

#### IV. DISCUSSION

The resistivity measurements suggest that antiferromagnetism in the high-pressure metallic state of  $\text{NiS}_2$  is

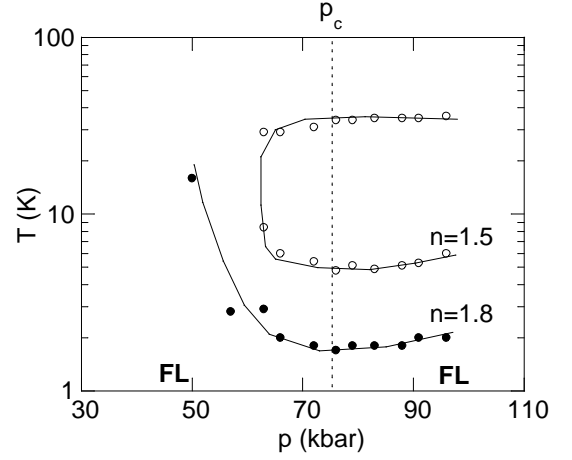


FIG. 7: Contours of the resistivity exponent  $n = \partial \ln \Delta\rho / \partial \ln T$  in the temperature-pressure plane for  $\text{NiS}_2$  in the metallic regime. A Fermi liquid (FL)  $T^2$  temperature dependence of the resistivity is seen at the lowest temperatures up to  $T_{FL}$  defined, somewhat arbitrarily, by the condition  $n(T_{FL}) = 1.8$ .  $T_{FL}$  reaches a minimum at  $p_c \approx 76$  kbar. A non-Fermi liquid  $T^n$  form of the resistivity with  $n \approx 3/2$  is seen over approximately a decade in temperature above a few K near  $p_c$ .

suppressed at a critical pressure of  $p_c = 76 \pm 5$  kbar. At  $p_c$ , we find that the  $T^2$  coefficient of the resistivity  $A$  has its maximum and the Fermi liquid crossover temperature  $T_{FL}$  defined in the caption of Figure 7 has its minimum. Above  $T_{FL}$  near  $p_c$  the resistivity has an approximately  $T^{3/2}$  temperature dependence over a decade in temperature. As already stated, this is the behaviour expected in the SCR model for a metal in 3D on the border of antiferromagnetism at low temperatures. Also, we note that  $p_c$  is of the same order of magnitude as that inferred for the correspondence between the temperature-composition and temperature-pressure phase diagram as discussed in the previous section.

The weakness or absence of a signature of  $T_N$  in the resistivity below  $p_c$  is not necessarily surprising since the same is the case for  $\text{NiS}_{2-x}\text{Se}_x$  in the range  $0.4 \leq x \leq 1$  (Refs. 12,33) where antiferromagnetic order has been confirmed by neutron scattering.<sup>10,34,35</sup> The signature of  $T_N$  in  $\rho$  can be washed out by inhomogeneities in composition or pressure and may only show up in high-precision measurements of  $\partial\rho/\partial T$  in the pressure range where  $T_N$  is not too strongly varying with pressure and where pressure is hydrostatic.

The fact that  $T_{FL}$  and correspondingly the  $T^2$  coefficient of the resistivity  $A$  remain finite at  $p_c$  is not necessarily inconsistent with our assumption that  $p_c$  marks the border of antiferromagnetism. The antiferromagnetic transition vs pressure may be first order as is found in a number of antiferromagnetic metals such as  $\text{YMn}_2$  and  $\text{GdMn}_2$ .<sup>36</sup> First order quantum phase transitions also seem to be common in ferromagnetic metals such as  $\text{MnSi}$  and  $\text{Ni}_3\text{Al}$ .<sup>5,37,38</sup>  $T_{FL}$  and  $A$  remain finite at the anti-

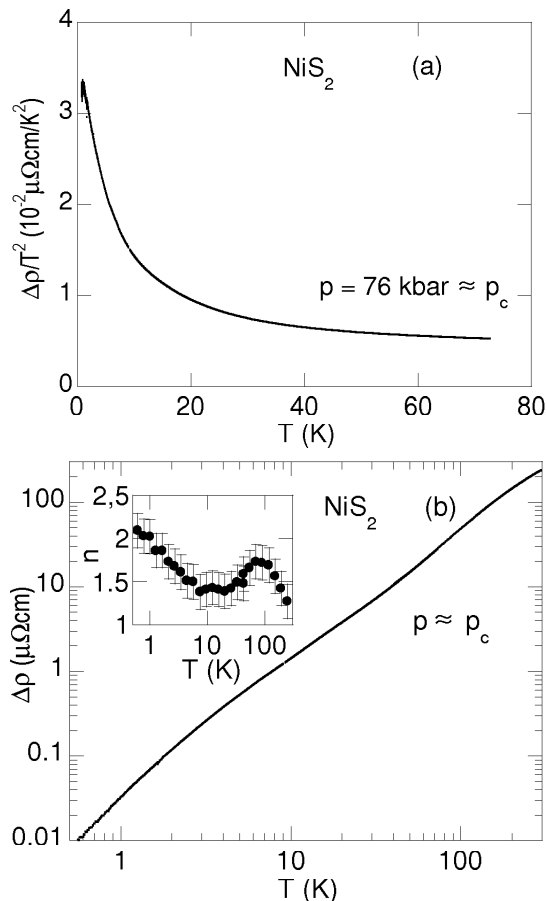


FIG. 8: The non-Fermi liquid form of the resistivity of NiS<sub>2</sub> near  $p_c \approx 76$  kbar. (a)  $\Delta\rho/T^2$  vs  $T$  does not saturate and thus has a non-Fermi liquid form over a wide temperature range, except at very low temperatures (Fig. 5). (b) A plot of  $\ln \Delta\rho$  vs  $\ln T$  and the resistivity exponent  $n = \partial \ln \Delta\rho / \partial \ln T$  vs  $\ln T$  exhibits an undulating form in contrast to the simple  $T^{3/2}$  temperature dependence seen in NiSSe (Ref. 12 and Fig. 1). This undulating form of the resistivity has been seen in different NiS<sub>2</sub> specimens and with two independent measurement systems.

ferromagnetic boundary of YMn<sub>2</sub>, a system that shares some features in common with NiS<sub>2</sub>. (We note, however, that the absence of Fermi liquid behaviour does not necessarily mean that the quantum phase transition is continuous.<sup>13,14</sup>)

The Fermi liquid crossover temperature,  $T_{FL}$ , in NiS<sub>2</sub> is very much lower than the characteristic spin fluctuation temperature,  $T_{sf}$ , which may be expected to be of the order of  $10^3$  K in a typical d-metal. We consider to what extent the behaviour above  $T_{FL}$ , but below  $T_{sf}$ , may be understood in terms of an itinerant-electron model for a continuous antiferromagnetic quantum critical point. The predictions of the SCR model as analyzed by Rosch has already been outlined in the introduction for an antiferromagnetic quantum critical point. With increasing reduced temperature  $t = T/T_{sf}$ ,

the SCR model predicts that the resistivity exponent  $n = \partial \ln \Delta\rho / \partial \ln T$  drops from an initial value of  $3/2$  towards unity around  $t \approx 1/(k_F l)$ , back to a value of order  $3/2$  around  $t \approx 1/\sqrt{k_F l}$  and then to zero for  $t \gg 1/(k_F l)$ .<sup>18,19,20</sup> This behaviour is qualitatively similar to that seen in NiS<sub>2</sub> at  $p_c$  and above  $T_{FL}$  (inset Fig. 8). For reasonable choices of parameters for NiS<sub>2</sub>,  $T_{sf} \approx 10^3$  K,  $k_F \approx 0.5 \text{ \AA}^{-1}$  and  $l \approx 140 \text{ \AA}^{-1}$ ,<sup>39</sup> we expect the minimum of  $n$  to occur near 15 K and the maximum at around 120 K. These crossover temperatures are in rough agreement with our observations.

In NiSSe,  $l$  is an order of magnitude smaller than in NiS<sub>2</sub> and thus the minimum and maximum of  $n$  are expected to arise at around 100 K and 320 K, respectively. Over the temperature range of the experiments shown in the inset of Figure 1 the model predicts a simple  $T^{3/2}$  temperature dependence without modulation, as is seen. The difference in the behaviour of NiS<sub>2</sub> and NiSSe at their respective critical conditions is thus not surprising. The combined effects of scattering from spin and lattice fluctuations might also lead to an undulating behaviour of  $n$ , but this would naively be expected to arise in both NiS<sub>2</sub> and NiSSe, in contradiction with observation.

## V. CONCLUSIONS

The temperature dependence of the resistivity of NiS<sub>2</sub> near the critical pressure, which has been found to be  $p_c = 76 \pm 5$  kbar, is consistent with that expected for a metal on the border of itinerant-electron antiferromagnetism. The Fermi liquid crossover temperature,  $T_{FL}$ , which defines the range over which the resistivity is roughly quadratic in temperature, does not vanish at  $p_c$ , but is three orders of magnitude smaller than the characteristic spin fluctuation temperature  $T_{sf}$ . The finite value of  $T_{FL}$  may indicate that the antiferromagnetic quantum critical point is first order, as is the case in related materials such as YMn<sub>2</sub>.

Over a wide temperature range above  $T_{FL}$  the temperature variation of the resistivity exhibits a non-Fermi liquid form that can be understood in terms of the effects of spin-fluctuation scattering at cold and hot spots of the Fermi surface as anticipated by Rosch in his refined treatment of the self-consistent-renormalization (SCR) model. In particular, the resistivity exponent  $n = \partial \ln \Delta\rho / \partial \ln T$  exhibits an undulating structure which is consistent not only qualitatively, but also approximately quantitatively with the predictions of this model. The model also accounts for the absence of this undulating structure in the related, but less pure material, NiSSe, which is known to be at the border of antiferromagnetism at ambient pressure.

## Acknowledgments

We thank A. Rosch and S. Julian for valuable discussions. PGN is grateful for support by the FERLIN pro-

gram of the European Science Foundation.

- 
- \* e-mail: philipp.niklowitz@frm2.tum.de; present address: Physik Department E21, Technische Universität München, 85747 Garching, Germany
- <sup>1</sup> J. A. Hertz, Phys.Rev.B **14**, 1165 (1976), and references therein.
  - <sup>2</sup> A. J. Millis, Phys.Rev.B **48**, 7183 (1993), and references therein.
  - <sup>3</sup> T. Moriya, *Spin Fluctuations in Itinerant Electron Magnetism* (Springer, Berlin, 1985), and references therein.
  - <sup>4</sup> G. G. Lonzarich, in *Electron*, edited by M. Springford (Cambridge University Press, Cambridge, 1997), chap. 6, pp. 109–147.
  - <sup>5</sup> P. G. Niklowitz, F. Beckers, G. G. Lonzarich, G. Knebel, B. Salce, J. Thomasson, N. Bernhoeft, D. Braithwaite, and J. Flouquet, Phys.Rev.B **72**, 24424 (2005).
  - <sup>6</sup> J. Mathon, Proc.Roy.Soc.A **306**, 355 (1968).
  - <sup>7</sup> C. M. Varma, P. B. Littlewood, S. Schmitt-Rink, E. Abrahams, and A. E. Ruckenstein, Phys.Rev.Lett. **63**, 1996 (1989).
  - <sup>8</sup> T. Holstein, R. E. Norton, and P. Pincus, Phys.Rev.B **8**, 2647 (1973).
  - <sup>9</sup> G. Baym and C. Pethick, *Landau-Fermi liquid theory* (Wiley, New York, 1991), chap. 3.
  - <sup>10</sup> M. Matsuura, H. Hiraka, K. Yamada, and Y. Endoh, J.Phys.Soc.Jpn. **69**, 1503 (2000).
  - <sup>11</sup> J. M. Honig and J. Spalek, Chem.Mater. **10**, 2910 (1998).
  - <sup>12</sup> S. Miyasaka, H. Takagi, Y. Sekine, H. Takahashi, N. Mori, and R. J. Cava, J.Phys.Soc.Jpn. **69**, 3166 (2000).
  - <sup>13</sup> C. Pfleiderer, S. R. Julian, and G. G. Lonzarich, Nature **414**, 427 (2001).
  - <sup>14</sup> N. Doiron-Leyraud, I. R. Walker, L. Taillefer, M. J. Steiner, S. R. Julian, and G. G. Lonzarich, Nature **425**, 595 (2003).
  - <sup>15</sup> D. Belitz, T. R. Kirkpatrick, and T. Vojta, Phys.Rev.Lett. **82**, 4707 (1999).
  - <sup>16</sup> P. Coleman, C. Pépin, Q. Si, and R. Ramazashvili, J.Phys.:Condens.Mat. **13**, R723 (2001).
  - <sup>17</sup> Q. Si, S. Rabello, K. Ingersent, and J. L. Smith, Nature **413**, 804 (2001).
  - <sup>18</sup> R. Hlubina and T. M. Rice, Phys.Rev.B **51**, 9253 (1995).
  - <sup>19</sup> A. Rosch, Phys.Rev.Lett. **82**, 4280 (1999).
  - <sup>20</sup> A. Rosch, Phys.Rev.B **62**, 4945 (2000).
  - <sup>21</sup> S. R. Julian, F. V. Carter, F. M. Grosche, R. K. W. Haselwimmer, S. J. Lister, N. D. Mathur, G. J. McMullan, C. Pfleiderer, S. S. Saxena, I. R. Walker, et al., J.Magn.Magn.Mat. **177-181**, 265 (1998).
  - <sup>22</sup> F. M. Grosche, M. J. Steiner, P. Agarwal, I. R. Walker, D. M. Freye, S. R. Julian, and G. G. Lonzarich, Physica B (2000).
  - <sup>23</sup> D. W. Bullett, J.Phys.C **15**, 6163 (1982).
  - <sup>24</sup> A. Y. Matsuura, H. Watanabe, C. Kim, S. Doniach, Z.-X. Shen, T. Thio, and J. W. Bennett, Phys.Rev.B **58**, 3690 (1998).
  - <sup>25</sup> J. A. Wilson and G. D. Pitt, Philosoph.Mag. **23**, 1297 (1971).
  - <sup>26</sup> Y. Sekine, H. Takahashi, N. Mori, T. Matsumoto, and T. Kosaka, Physica B **237-238**, 148 (1997).
  - <sup>27</sup> J. A. Wilson, Adv. in Phys. **21**, 143 (1972).
  - <sup>28</sup> J. A. Wilson, in *The Metallic and Non-Metallic States of Matter*, edited by P. P. Edwards and C. N. R. Rao (Taylor and Francis, London, 1997), chap. 9, pp. 215–260.
  - <sup>29</sup> N. Mori and H. Takahashi, J.Magn.Magn.Mat. **31-34**, 335 (1983).
  - <sup>30</sup> A. Husmann, J. Brooke, T. F. Rosenbaum, X. Yao, and J. M. Honig, Phys.Rev.Lett. **84**, 2465 (2000).
  - <sup>31</sup> J. Wittig, Z.Phys. **195**, 215 (1966).
  - <sup>32</sup> A. Eichler and J. Wittig, Z.Angew.Phys. **25**, 319 (1968).
  - <sup>33</sup> X. Yao, J. M. Honig, T. Hogan, C. Kannewurf, and J. Spalek, Phys.Rev.B **54**, 17469 (1996).
  - <sup>34</sup> S. Sudo and T. Miyadai, J.Phys.Soc.Jpn. **54**, 3934 (1985).
  - <sup>35</sup> S. Sudo, J.Magn.Magn.Mat. **114**, 57 (1992).
  - <sup>36</sup> R. Hauser, A. Indinger, E. Bauer, and E. Gratz, J.Magn.Magn.Mat. **140-144**, 799 (1995).
  - <sup>37</sup> C. Pfleiderer, G. J. McMullan, S. R. Julian, and G. G. Lonzarich, Phys.Rev.B **55**, 8330 (1997).
  - <sup>38</sup> C. Thessieu, C. Pfleiderer, A. N. Stepanov, and J. Flouquet, J.Phys.Cond.Mat. **9**, 6677 (1997).
  - <sup>39</sup> The order of magnitude,  $T_{sf}$ , was estimated from the value of  $A$  in the SCR model,  $k_F$  from the size of the Brillouin zone and electron filling of the bands and  $l$  from  $\rho_0$ .